

# MATERIAL VERIFICATION REPORT

Job No : 18056

Project : JALAN

: JALAN TOL JAPEK2 ELEVATED LONGSPAN GRAND WISATA

Client: PT. WASKITA KARYA

PO No. :

STRUCTURAL

Durbe

: 19 Jan 2019

Page

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Material Type

: BOLTING

Report No

1 WESE-PTG-STRANG-007

Stone Man.

2 00

No	Description Material / Specification / Size	Otty	Class	Identification			Mill Cert	Witnessell		MICIRI	TO			
				Heat No	Product No	No		Surf. Cond.	Straig.	No.	Mo.	Status		
												A.	PH.	1
	STUD BOLT / ISO 13918 TYPE SD1 / Catton Steel Diameter 25mm X 250mm Length	34804		3/3 NBOT 29	PHE HEA		Attach	Good	Glood			X.		H
2	PERFIUL CERAMIC / IGO 13918 Type UF 25 / Diameter 25 mm	34804		MIA			Attach	Good	Good			ж	Ħ	F

Remarks: Stud Bolt for production and PER

Prepared by.

Approved by.

PT Gunanusa Utama Fabricators KSO WASKITA- Acted VIRAMA KARYA

JIC (\*) IF ANY SINCONFORMITIES AND SEXTEND FOR ANY OBSERVATION REPORT OF METHOD WILL BE ATTACHED

Legend: A = Accept, H = Hold, R = Reject

# Raw Material Verification Quality Manual Template

Harish K. Sharma, Nicolas Y.
Njintang, Rekha S. Singhal, Pragati
Kaushal

#### **Raw Material Verification Quality Manual Template:**

FDA Inspection Operations Manual .1990 Pharmaceutical Packaging Technology D. A. Dean, E. R. Evans, I. H. Hall, 2005-07-12 Pharmaceutical packaging requires a greater knowledge of materials and a greater intensity of testing than most other packed products not to mention a sound knowledge of pharmaceutical products and an understanding of regulatory requirements Structured to meet the needs of the global market this volume provides an assessment of a wide range of issues It covers the entire supply chain from conversion of raw materials into packaging materials and then assembled into product packs Integrating information from many drug delivery systems the author discusses testing and evaluation and emphasizes traceability and the need to for additional safeguards **Tool and Manufacturing Engineers** Handbook: Plastic Part Manufacturing Philip Mitchell, 1996-12-09 This volume focuses on the practical application of processes for manufacturing plastic products It includes information on design for manufacturability DFM material selection process selection dies molds and tooling extrusion injection molding blow molding thermoforming lamination rotational molding casting foam processing compression and transfer molding fiber reinforced processing assembly and fabrication quality plant engineering and maintenance management The Army Lawyer ,1991 **Food Industries Manual** Christopher G.J. Baker, M.D. Ranken, R.C. Kill, 2012-12-06 It is a measure of the rapidity of the changes The work has been revised and updated and taking place in the food industry that yet another following the logic of the flow sheets there is some edition of the Food Industries Manual is required simplification and rearrangement among the chap after a relatively short interval As before it is a ters Food Packaging now merits a separate pleasure to be involved in the work and we hope chapter and some previous sections dealing mainly that the results will continue to be of value to with storage have been expanded into a new readers wanting to know what how and why the chapter covering Food Factory Design and Opera food industry does the things which it does tions For this edition we have made a major depar There is one completely new chapter entitled ture from the style of earlier editions by comple Alcoholic Beverages divided into Wines Beers tely revising the layout of many of the chapters and Spirits There is a strain of thought which Previously the chapters were arranged as a series does not yet consider the production of those of notes on specific topics set out in alphabetical drinks to be a legitimate part of the food industry order in the manner of an encyclopaedia Agriculture Handbook ,1981 Set includes revised editions of some Setting Up a Quality Control System United States. Small Business Administration. Management Assistance issues Division. Support Services Section, 1982 **Boating** ,1972-07 The Certified Pharmaceutical GMP Professional **Handbook, Second Edition** Mark Allen Durivage, 2016-05-26 The purpose of this handbook is to assist individuals for the Certified Pharmaceutical Good Manufacturing Practices Professional CPGP examination and provide a reference for the practitioner The second edition reflects the Body of Knowledge which was updated in 2015 This edition has also incorporated additional information including updated references. The updates reflect the current trends and expectations of the evolving

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Food Industry Quality Control Systems Mark Clute, 2008-10-22 With more than 100 tables spreadsheets and program outlines Food Industry Quality Control Systems features all of the systems needed for food manufacturing facilities to assure food quality and safety This book outlines the role of management in overall quality control and the role of HACCP within a company It explains the basics of quality control programs including pest control programs lot coding organizational charts specifications supplier certification and sanitation programs Readers can visit the CRC website to download supporting programs in a plug and play format and adaptable forms with specific examples and resources that enable companies to customize forms to fit their own needs Iso 9001 Arpad Gaal, 2001-06-28 ISO 9001 2000 for Small Business Management Implementing Process Approach Quality Management demonstrates how a process approach quality management system performs in the real work environment The book gives you an ISO based quality management tool featuring the year 2000 requirements for ISO 9001 It includes the quality system manual the oper The Certified Manager of Ouality/Organizational Excellence Handbook, Fourth Edition Russell T. Westcott, 2013-10-07 This handbook is a comprehensive reference source designed to help professionals address organizational issues from the application of the basic principles of management to the development of strategies needed to deal with the technological and societal concerns of the new millennium The content of this fourth edition has been revised to reflect a more current global perspective and to match the updated Body of Knowledge BoK of ASQ s Certified Manager of Quality Organizational Excellence CMQ OE In order to provide a broad perspective of quality management this book has specifically been written to address Historical perspectives relating to the evolution of particular aspects of quality management including recognized experts and their contributions Key principles concepts and terminology relevant in providing quality leadership and communicating quality needs and results Benefits associated with the application of key concepts and quality management principles Best practices describing recognized approaches for good quality management Barriers to success including common problems that the quality manager might experience when designing and implementing quality management and insights as to why some quality initiatives fail Guidance for preparation to take the CMQ OE examination Organized to follow the BoK exactly throughout each section of this handbook the categorical BoK requirements associated with good quality management practices for that section are shown in a box preceding the pertinent text These BoK requirements represent the range of content and the cognitive level to which multiple choice questions can be presented Although this handbook thoroughly prepares individuals for the ASQ CMQ OE exam the real value resides in post exam usage as a day to day reference source

for assessing quality applications and methodologies in daily processes. The content is written from the perspective of practitioners and its relevance extends beyond traditional product quality applications Handbook of Validation in Pharmaceutical Processes, Fourth Edition James Agalloco, Phil DeSantis, Anthony Grilli, Anthony Pavell, 2021-10-28 Revised to reflect significant advances in pharmaceutical production and regulatory expectations Handbook of Validation in Pharmaceutical Processes Fourth Edition examines and blueprints every step of the validation process needed to remain compliant and competitive This book blends the use of theoretical knowledge with recent technological advancements to achieve applied practical solutions As the industry's leading source for validation of sterile pharmaceutical processes for more than 10 years this greatly expanded work is a comprehensive analysis of all the fundamental elements of pharmaceutical and bio pharmaceutical production processes Handbook of Validation in Pharmaceutical Processes Fourth Edition is essential for all global health care manufacturers and pharmaceutical industry professionals Key Features Provides an in depth discussion of recent advances in sterilization Identifies obstacles that may be encountered at any stage of the validation program and suggests the newest and most advanced solutions Explores distinctive and specific process steps and identifies critical process control points to reach acceptable results New chapters include disposable systems combination products nano technology rapid microbial methods contamination control in non sterile products liquid chemical sterilization and medical device manufacture **Operations Management** David Barnes, 2018-01-30 This fascinating new core textbook authored by a highly respected academic with over a decade of industry experience takes a global and strategic approach to the important topic of operations management OM Integrating contemporary and traditional theories the text covers everything a student needs to understand the reality of operations in the modern world and combines the latest cutting edge thinking with innovative learning features Written in a concise and engaging style and based on up to date research in the field the book provides a range of international case studies and examples that help students to apply theoretical knowledge to real world practice This is a must have textbook for students studying operations management modules on undergraduate postgraduate and MBA programmes In addition this is an ideal textbook to accompany modules on operations strategy production management and services management Heat Exchanger Design Handbook, Second Edition Kuppan Thulukkanam, 2013-05-20 Completely revised and updated to reflect current advances in heat exchanger technology Heat Exchanger Design Handbook Second Edition includes enhanced figures and thermal effectiveness charts tables new chapter and additional topics all while keeping the qualities that made the first edition a centerpiece of information for practicing engineers research engineers academicians designers and manufacturers involved in heat exchange between two or more fluids See What s New in the Second Edition Updated information on pressure vessel codes manufacturer s association standards A new chapter on heat exchanger installation operation and maintenance practices Classification chapter now includes coverage of scrapped surface graphite coil wound microscale and printed circuit heat exchangers Thorough revision

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